

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Applicant: Malcolm Wilson MOON et al.
Title: Mannich Base Prodrugs of 3-(Pyrrol-2-yl-methylidene)-2-Indolinone Derivatives
Appl. No.: Unassigned
Filing Date: 12/24/2003
Examiner: Unassigned
Art Unit: Unassigned

INFORMATION DISCLOSURE STATEMENT
UNDER 37 CFR §1.56

Mail Stop PATENT APPLICATION
Commissioner for Patents
PO Box 1450
Alexandria, Virginia 22313-1450

Sir:

Applicants submit herewith on Form PTO/SB/08 a listing of the documents cited by or submitted to the U.S. PTO in parent application Serial No. 09/863,804, filed 05/24/2001. As provided in 37 CFR §1.98(d), copies of the documents are not being provided since they were previously submitted to the United States Patent & Trademark Office in the above-identified parent application.

The submission of any document herewith, which is not a statutory bar, is not intended as an admission that such document constitutes prior art against the claims of the present application or that such document is considered material to patentability as defined in 37 CFR §1.56(b). Applicants do not waive any rights to take any action which would be appropriate to antedate or otherwise remove as a competent reference any document which is determined to be a *prima facie* art reference against the claims of the present application.

TIMING OF THE DISCLOSURE

The listed documents are being submitted in compliance with 37 CFR §1.97(b), within three (3) months of the filing date of the application.

RELEVANCE OF EACH DOCUMENT

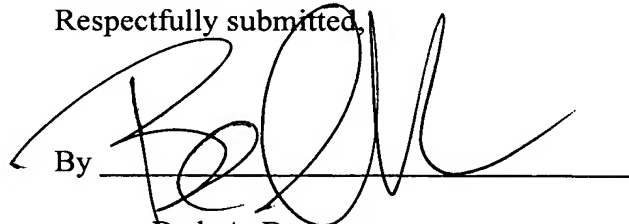
The relevance of the foreign-language documents is explained in the parent application.

Applicants respectfully request that any listed document be considered by the Examiner and be made of record in the present application and that an initialed copy of Form PTO/SB/08 be returned in accordance with MPEP §609.

The Commissioner is hereby authorized to charge any additional fees which may be required regarding this application under 37 CFR §§ 1.16-1.17, or credit any overpayment, to Deposit Account No. 19-0741. Should no proper payment be enclosed herewith, as by a check being in the wrong amount, unsigned, post-dated, otherwise improper or informal or even entirely missing, the Commissioner is authorized to charge the unpaid amount to Deposit Account No. 19-0741.

Respectfully submitted,

By

A handwritten signature in black ink, appearing to read 'Beth A. Burrous', is written over a horizontal line.

Beth A. Burrous
Attorney for Applicant
Registration No. 35,087

Date: December 24, 2003

FOLEY & LARDNER

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Substitute for form 1449B/PTO INFORMATION DISCLOSURE STATEMENT BY APPLICANT Date Submitted: December 24, 2003 <i>(use as many sheets as necessary)</i>				C mplete if Known	
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				First Named Inventor	Malcolm Wilson MOON
				Group Art Unit	Unassigned
				Examiner Name	Unassigned
				Attorney Docket Number	034536-0920
Sheet	1	of	11		

U.S. PATENT DOCUMENTS						
Examiner Initials*	Cite No. ¹	U.S. Patent Document		Name of Patentee or Applicant of Cited Document	Date of Publication of Cited Document MM-DD-YYYY	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear
		Number	Kind Code ² (if known)			
	A1	2968557		Burgandt et al.	01/17/1961	
	A2	4002749		Rovnyak	01/11/1977	
	A3	4053613		Rovnyak et al.	10/11/1977	
	A4	4642309		Michel et al.	02/10/1987	
	A5	4826847		Michel et al.	05/02/1989	
	A6	5051417		Nadler et al.	09/24/1991	
	A7	5124347		Connor et al.	06/23/1992	
	A8	5196446		Levitzki et al.	03/23/1993	
	A9	5302606		Spada et al.	04/12/1994	
	A10	5322950		Sircar et al.	06/21/1994	
	A11	5374652		Buzzetti et al.	12/20/1994	
	A12	5382593		Le Baut et al.	01/17/1995	
	A13	5389661		Sircar et al.	02/14/1995	
	A14	5397787		Buzzetti et al.	03/14/1995	
	A15	5409949		Buzzetti et al.	04/25/1995	
	A16	5792783		Tang et al.	08/11/1998	
	A17	5834504		Tang et al.	11/10/1998	
	A18	5849710		Battistini et al.	12/15/1998	
	A19	5880141		Tang et al.	03/9/1999	

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		U.S. Patent Document			
	A20	5883113		Tang et al.	03/16/1999
	A21	5883116		Tang et al.	03/16/1999
	A22	5886020		Tang et al.	03/23/1999
	A23	6133305		Tang et al.	10/17/2000
	A24	6451838		Moon et al.	09/2002

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	A25	WO	91/13055			09/05/1991		
	A26	WO	92/07830			05/14/1992		
	A27	WO	92/20642			11/26/1992		
	A28	WO	93/01182			01/21/1993		
	A29	WO	94/14808			07/07/1994		
	A30	WO	95/01349			01/12/1995		
	A31	WO	95/17181			06/29/1995		
	A32	WO	96/00226			01/04/1996		
	A33	WO	96/16964			06/06/1996		
	A34	WO	96/22976			08/01/1995		
	A35	WO	96/32380			10/17/1996		
	A36	WO	96/40116			12/19/1996		

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	A37	WO	97/25986			07/24/1997		
	A38	WO	98/07695			02/26/98		
	A39	WO	98/24432			6/11/98		
	A40	WO	98/38984			9/11/98		
	A41	WO	98/50356			11/12/98		
	A42	WO	99/10325			3/4/99		
	A43	WO	99/52869			10/21/99		
	A44	WO	99/61422			12/2/99		
	A45	WO	99/65869			12/23/99		
	A46	WO	00/08202			2/17/00		
	A47	WO	00/35906			6/22/00		
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	A52	WO	01/60814			8/23/01		
	A53	WO	01/90104			11/29/01		
	A54	DE	2159360			11/30/71		Trans.
	A55	DE	2159361			11/30/71		Trans.

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	A56	DE	2159362			11/30/71		
	A57	DE	2159363			11/29/71		Trans.
	A58	DE	2321656			4/28/73		Trans.
	A59	DE	3426419			1/23/86		Trans.
	A60	EP	0252713			9/12/1990		
	A61	EP	0351213			1/17/90		
	A62	EP	0525472			2/3/93		
	A63	EP	0632102			1/4/95		Trans.
	A64	EP	0662473			7/12/95		
	A65	EP	0788890			8/13/97		
	A66	EP	0769947			5/2/97		
	A67	EP	0934931			8/11/99		
	A68	EP	1082305	A1		3/14/01		
	A69	FR	1398224			3/29/65		Trans.
	A70	FR	1599772			8/1970		Trans.
	A71	FR	2689397			10/8/93		Trans.
	A72	FR	809691			3/4/59		
	A73	FR	835473			5/18/60		
	A74	JP	6229570			7/2/87		Trans.

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	A75	JP	6239564			2/20/87		Trans.
	A76	JP	63141955			6/14/88		Trans.
	A77	JP	558894			3/9/93		Trans.
	A78	CA	2012634			9/20/91		
	A79	AU	286870			5/11/67		

NON PATENT LITERATURE DOCUMENTS			
Examiner Initials*	Cite No. ¹	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.) date, page(s), volume-issue number(s), publisher, city and/or country where published.	T ⁶
	A80	Andreani et al., "Potential Antitumor Agents. 25[1]. Synthesis and Cytotoxic Activity of 3-(2-Chloro-3-Indolymethylene)1,3-Dihydroindol-2-Ones," <u>Anticancer Research</u> 16:3585-3588 (1996) 8 Elsevier, Paris	
	A81	Andreani et al., "Synthesis and cardiotoxic activity of 2-indolinones," <u>Eur. J. Med. Chem.</u> 25:187-190 (1990)	
	A82	Andreani et al., "Synthesis and cardiotoxic activity of 2-indolinones bearing pyridyl groups," <u>Eur. J. Med. Chem.</u> 28:653-657 (1993) 8 Elsevier, Paris	
	A83	Andreani et al., "Synthesis and cardiotoxic activity of pyridylmethylene-2-indolinones," <u>Eur. J. Med. Chem.</u> 27:167-170 (1992) 8 Elsevier, Paris	
	A84	Andreani et al., "Synthesis and potential coanthracyclenic activity of substituted 3-(5-imidazo[2,1-b]thiazolylmethylene)-2-indolinones," <u>Eur. J. Med. Chem.</u> 32:919-924 (1997)8 Elsevier, Paris	
	A85	Andreani et al., "Synthesis of lactams with potential cardiotoxic activity," <u>Eur. J. Med. Chem.</u> 28:825-829 (1993)	
	A86	Andreani et al., "In Vivo Cardiotoxic Activity of Pyridylmethylene-2-indolinones," <u>Arzneimittel-Forschung Drug Research</u> 48:727-729 (1998) 8	
	A87	Bahner and Brotherton, "9-(4-Aminobenzylidene)fluorenes," <u>J. Med. Chem.</u> 12:722-723 (1969)	

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	A88	Bahner et al., "Benzylideneindenes with Oxygen Attached to the Indene Ring," <u>J. Med. Chem.</u> 12:721-722 (1969)	
	A89	Bamfield et al., "Diels-Alder Reactions of Oxindolylideneacetone," <u>J. Chem. Soc. (C)</u> 1028-1030 (1966) 8	
	A90	Borsche et al., "Über vielkernige kondensierte Systeme mit heterocyclischen Ringen. XIII.," <u>Liebigs Ann. Chem.</u> 550:160-174 (1941)	
	A91	Buzzetti et al., "Cinnamamide Analogs as Inhibitors of Protein Tyrosine Kinases," <u>Il Farmaco</u> 48:615-636 (1993)	
	A92	Chatten et al., "Substituted Oxindoles. Part VI. Polarographic Reduction of Substituted <i>trans</i> -3-Benzylideneindol-2(3 <i>H</i>)-ones," <u>J. Chem. Soc. Perkin II</u> : 469-473 (1973)	
	A93	Coda et al., "(Z)- and (E)-Arylidene-1,3-dihydroindol-2-ones: Configuration, Conformation and Infrared Carbonyl Stretching Frequencies," <u>J. Chem. Soc. Perkin Trans. II</u> : 615-619 (1984)	
	A94	Coda et al., "3-(4-methylbenzylidene)-1,3-dihydroindol-2-one," <u>Journal of the Chemical Society, Perkin Transactions 2</u> 4:615-620 (1984) DATABASE CROSSFIRE, Beilstein Reference No. 6-21	
	A95	Decodts et al., "Suicide inhibitors of proteases. Lack of activity of halomethyl derivatives of some aromatic lactams," <u>Eur. J. Med. Chem.</u> 18: 107-111 (1983)	
	A96	Desimoni et al., "Catalysis with Inorganic Cations. V ¹ Intramolecular Hetero Diels-Alder <i>versus</i> Ene Reactions: Effect of Magnesium perchlorate on Chemoselectivity," <u>Tetrahedron</u> 52(36) 12009-12018 (1196) 8 Pergamon	

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	A97	Elliott and Rivers, "Reduction of Some Oxindolylidene Derivatives to 3-Substituted Oxindoles by Sodium Borohydride," <u>J. Med. Chem.</u> 29:2438-2440 (1964)	
	A98	Elliott et al., "1-methyl-2-(3-oxindolidenmethyl)-pyridinium," <u>Journal of Organic Chemistry</u> 29:2438-2440 (1964) DATABASE CROSSFIRE, Beilstein Reference No. 5-24	
	A99	Gazit et al., "Tyrophostins. 2. Heterocyclic and α -Substituted Benzylidenemalononitrile Tyrophostins as Potent Inhibitors of EGF Receptor and ErbB2/neu Tyrosine Kinases," <u>J. Med. Chem.</u> 34:1896-1907 (1991) copyright Am. Clem. Soc.	
	A100	Hirao et al., "Rhodium-Catalyzed Carbonylation of 2-Alkynylaniline: Syntheses of 1,3-Dihydroindol-2-ones," <u>Tetrahedron Letters</u> 36(35) 1995 8Pergamon	
	A101	Hodges et al., "Chemical and biological properties of some oxindolylid-3-methines," <u>Canadian J. Chemistry</u> 46:2189-2194 (1968)	
	A102	Howard, Harry R., "Lactam Derivatives," U.S. Provisional Patent Application Number 60/015134	
	A103	Howard et al., "Synthesis and aldose reductase inhibitory activity of substituted 2(1H)-benzimidazolone- and oxindole-1-acetic acids," <u>Eur. J. Med. Chem.</u> 27:779-789 (1992) 8 Elsevier, Paris	
	A104	Katritzky et al., "Color and Constitution. Part 8[1]. Some Novel Dyestuffs Containing Indoxyl Residues," <u>J. Heterocyclic Chem.</u> 25:1287-1292 (1988)	
	A105	Kobayashi et al., "Anti-tumor Activity of Indole Derivatives," <u>Yakugaku Zasshi</u> 97:1033-1039 (1977)	

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		First Named Inventor	Malcolm Wilson MOON
		Group Art Unit	Unassigned
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		Attorney Docket Number	034536-0920

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	A106	Kovac and Stetinova, "Furan derivatives. LXXX. Synthesis and properties of substituted furfurylidenoxindoles," <u>Chem. rvesu</u> 30:484-492 (1976)	
	A107	Levitzki and Gazit, "Tyrosine Kinase Inhibition: An Approach to Drug Development," <u>Science</u> 267:1782-1788 (1995)	
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	A109	Mohammadi et al., "Structures of the Tyrosine Kinase Domain of Fibroblast Growth Factor Receptor in Complex with Inhibitors," <u>Science</u> 276:955-960 (1997) 8 American Association for the Advancement of Science	
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	A111	Nodiff et al., "Antimalarial Phenanthrene Amino Alcohols. 3. Halogen-containing 9-phenanthrenemethanols," <u>Chemical Abstracts</u> , Vol. 83, abstract no. 188214 (1975)	
	A112	O'Sullivan and Rothery, "The Preparation and Possible Clinical Significance of 4'-Dialkylaminoisoidogenides," <u>Clinica Chimica Acta</u> 62:181-182 (1975) 8Elsevier Scientific Publishing Company	
	A113	Pavlenko et al., "Introduction of aminomethyl groups into heterocyclic CH-acid molecules," <u>Dopov. Akad. Nauk Ukr Rsr.</u> , Ser. B: <u>Geol., Khim. Biol. Nauki</u> 7:64-66 (1980)	
	A114	Plowman et al., "Receptor Tyrosine Kinases as Targets for Drug Intervention," <u>DN&P</u> 7:334-339 (1994)	

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	A115	Quallich et al., "A General Oxindole Synthesis," <u>J. Synthetic Organic Chemistry</u> : 51-51 (1993)	
	A116	Schuchter et al., "Successful Treatment of Murine Melanoma with Bryostatin 1," <u>Cancer Research</u> 51:682-687 (1991)	
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	A119	Singh et al., "Indolinone Derivatives as Potential Antimicrobial Agents," <u>Zentralbl. Mikrobiol.</u> 144:105-109 (1989) copyright VEB Gustav Fischer Verlag Jena	
	A120	Spada, et al., "Small molecule inhibitors of tyrosine kinase activity," <u>Expert Opinion on Therapeutic Patents</u> 5:805-817 (1995) 8Ashley Publications	
	A121	Sun et al., "Design, Synthesis, and Evaluations of Substituted 3-[(3- or 4-Carboxyethylpyrrol-2-yl)methylidene]indolin-2-ones as Inhibitors of VEGF, FGF, and PDGF Receptor Tyrosine Kinases," <u>Journal of Medicinal Chemistry</u> 42: 5120-5130 (1999) 8American Chemical Society	
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	A123	Tacconi and Marinone, "Preparazione e caratteristiche di alcuni 3-ossindolidenderivati," <u>Ricerca Scientifica</u> 38:1239-1244 (1968)	

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	A124	Tacconi et al., "(Z)- and (E)-3-Alkylidene-1,3-dihydroindol-2-ones: Influence of Configuration on the Transmission of the Inductive Effect to the Carbonyl Group," <u>J.C.S. Perkin II</u> 150-154 (1976)	
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	A126	Traxler, "Protein tyrosine kinase inhibitors in cancer treatment," <u>Expert Opinion on Therapeutic Patents</u> 7(6):571-588 (1997) 8 Ashley Publications Ltd.	
	A127	Wahl et al., "3-benzilidene-5-methyl-1,3-dihydroindol-2-one," <u>Ann. Chim.</u> 350 (1926), DATABASE CROSSFIRE, Beilstein Reference No. 2-21-00-00290	
	A128	Wahl, Beilstein Reg. No. 191439, <u>Bull. Soc. Chim. Fr.</u> , page 1038 (1909)	
	A129	Wahl, Beilstein Reg. No. 231732, <u>Bull. Soc. Chim. Fr.</u> , pages 1035-1038 (1909)	
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	A131	Wright et al., "Cyclic Hydroxamic Acids Derived from Indole," <u>J. Am. Chem. Soc.</u> 78:221-224 (1956)	
	A132	Wright et al., "Inhibition of Angiogenesis in Vitro and In Ovo With an Inhibitor of Cellular Protein Kinases, MDL 27032," <u>J. Cellular Physiology</u> 152:448-457 (1992)	

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	A134	Zhungietu et al., "Reaction of Indoles and 2-Ketoindolines With Some Aldehydes," <u>Chemical Abstracts</u> , Vol. 78, abstract no. 111201 (1990)		

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